What is claimed is:

1. A method for treatment of chronic pain comprising orally administering a combination of a low dose of a tricyclic antidepressant compound and a standard dose of a non-narcotic analgesic.

2. The method of claim 1 wherein said tricyclic antidepressant is administered in a dosage of from about 2.5 mg to about 25 mg daily.

3. The method of claim 1 wherein said tricyclic antidepressant compound is selected from the group consisting of doxepin, amitriptyline, desipramine, imipramine and physiologically acceptable acid addition salts thereof.

4. The method of claim 1 wherein said physiologically acceptable acid addition salts are selected from the group consisting of the hydrochloride, hydrobromide, hydroiodide, acetate, valerate and oleate.

5. The method of claim 1 wherein said non-narcotic analgesic is administered in a dosage from about 0.50 gms to about 2.6 gms daily.

6. The method of claim 1 wherein said non-narcotic analgesic is selected from the group consisting of acetaminophen and NSAIDs.

7. The method of claim 1 wherein said low dose of tricyclic antidepressant compound and said standard dose of non-narcotic analgesic are present in a single composition including a pharmaceutically acceptable vehicle for oral administration.

8. The method of claim 7 wherein said composition is in a form selected from the group consisting of tablets, capsules, caplets, oral solutions, and oral suspensions.

9. A composition for treatment of chronic pain comprising a combination of a low dose of a tricyclic antidepressant compound and a standard dose of a non-narcotic analgesic in a pharmaceutical acceptable vehicle for oral administration.

10. The composition of claim 9 wherein said tricyclic antidepressant compound is administered in a dosage of from about 2.5 mg to about 25 mg daily.

11. The composition of claim 9 wherein said tricyclic antidepressant compound is selected from the group consisting of doxepin, amitriptyline, desipramine, imipramine, and physiologically acceptable acid addition salts thereof.

12. The composition of claim 9 wherein said physiologically acceptable acid addition salts are selected from the group consisting of the hydrochloride, hydrobromide, hydroiodide, acetate, valerate and oleate.

13. The composition of claim 9 wherein said non-narcotic analgesic is administered in a dosage for from about 0.50 gms to about 2.6 gms daily.

14. The composition of claim 9 wherein said non-narcotic analgesic is selected from the group consisting of acetaminophen or NSAIDs.

15. The composition of claim 7 wherein the combination of a tricyclic antidepressant and a non-narcotic analgesic and a pharmaceutically acceptable vehicle is in a form selected

from the group consisting of tablets, capsules, caplets, oral solutions and oral suspensions.